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sealed with aluminum flip-off type caps. FVIIa may be produced as disclosed in EP patent No. 200,421.

CLAIMS

- A pharmaceutical composition for parenteral administration, which comprises a peptide and dimethyl sulfone.
 - 2. A pharmaceutical composition according to claim 1, wherein the amount of dimethyl sulfone is of from 40 to 400 mM.
 - 3. A pharmaceutical composition according to claim 2, wherein amount of dimethyl sulfone is of from 125 to 350 mM.
 - 4. A pharmaceutical composition according to claim 1, wherein the composition is a solution.
 - 5. A pharmaceutical composition according to claim 1, wherein the composition is a suspension.
- 6. A pharmaceutical composition according to claim 1, which is suitable for administration by injection or infusion.
 - 7. A pharmaceutical composition according to claim 6, which is suitable for subcutaneous administration.
- 8. A pharmaceutical composition according to claim 6, which is suitable for intramuscular administration.
 - 9. A pharmaceutical composition according to claim 6, which is suitable for intravenous administration.
 - 10. A pharmaceutical composition according to claim 1, which is suitable for pulmonal administration.
- 11. A pharmaceutical composition according to claim 1, which is suitable for ophthalmic administration or topical administration.

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- 12. A pharmaceutical composition according to claim 1, wherein the peptide is human growth hormone, GLP-1, GLP-2, insulin, Factor VIIa, Factor VIII, erythropoeitin (EPO), glucagon, interleukin, such as interleukin-2 (IL-2), interferon- α or interferon- β , or an analogue thereof, or a derivative of any such peptide or analogue.
- 13. A pharmaceutical composition according to claim 12, wherein the peptide is human insulin or an analogue thereof, or a derivative of human insulin or the human insulin analogue.
- 10 14. A pharmaceutical composition according to claim 13, wherein the peptide is human insulin.
 - 15. A pharmaceutical composition according to claim 13, wherein the peptide is Asp(B28)-human insulin.
 - 16. A pharmaceutical composition according to claim 13, wherein the peptide is Lys(B28) Pro(B29)-human insulin.
- 17. A pharmaceutical composition according to claim 13, wherein the peptide isLys(B3) Glu(B29)-human insulin.
 - 18. A pharmaceutical composition according to claim 13, wherein the peptide is N^{cB29}-tetra-decanoyl des (B30)-human insulin.
- 19. A pharmaceutical composition according to claim 13, wherein the peptide is Gly(A21) Arg(B31) Arg(B32)-human insulin.
 - 20. A pharmaceutical composition according to claim 13, wherein the peptide is N^{εB29}–lito-choloyl-γ-glutamyl des (B30)-human insulin.
 - 21. A pharmaceutical composition according to claim 12, wherein the peptide is Gly(8)-human GLP-1.
- 22. A pharmaceutical composition according to claim 12, wherein the peptide is Arg(34), N-ε (γ-Glu(N-α-hexadecanoyl))-Lys(26)-human GLP-1(7-37)OH.

23. A pharmaceutical composition according to claim 12, wherein the peptide is Gly(2)-human GLP-2.